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Substitute for form 1449B/PTO INFORMATION DISCLOSURE STATEMENT BY APPLICANT

Sheet

Date Submitted:

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	Complete if Known
Application Number	10/600 868
Filing Date	12/22/2000
First Named Inventor	Ken Lipson
Group Art Unit	1614
Examiner Name	SPIVACK
Attorney Docket Number	038602-0994

				U.S. PATENT DOCUMENTS	· .	Pages, Columns, Lines,
Examiner	Cite	U.S. Patent I	Kind	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Where Relevant Passages or Relevant Figures Appear
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Examiner Signature	Phullis Spivack	Date Considered	1/23/05
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<sup>\*</sup>EXAMINER: Initial if reference coasidered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

<sup>&</sup>lt;sup>1</sup> Unique citation designation number. <sup>2</sup>See attached Kinds of U.S. Patent Documents. <sup>3</sup>Enter Office that issued the document, by the two-letter code (WIPO Standard Onique citation designation number. See attached Kinds of U.S. Patent Documents. Enter Onice that issued the document, by the two-letter code (WIFO Stant St. 3). For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. Skind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. Applicant is to place a check mark here if English language Translation is attached.

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	Substitute for form 1449B/PTO				Complete if Known			
	INFORMATION DIS	SCLO	SURE	Application Number				
	STATEMENT BY APPLICANT			Filing Date	12/22/2000			
	Date Submitted:  (use as many sheets as necessary)			First Named Inventor	Ken Lipson			
				Group Art Unit	1614			
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Sheet ·	2	of	30	Attorney Docket Number	038602-0994			

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Examiner Initials*	Cite No.1	U.S. Patent Number	Nocument Kind Code² (if known)	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
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	Examiner Signature	Phullis	Sawack	Date Considered	1/23/05
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¹ Unique citation designation number. ²See attached Kinds of U.S. Patent Documents. ³Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). ⁴For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. ⁵Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. ⁴Applicant is to place a check mark here if English language Translation is attached.

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Substitute for form 1449B/PTO		Complete if Known
INFORMATION DISCLOSURE STATEMENT BY APPLICANT	Application Number Filing Date First Named Inventor	12/22/2000 Ken Lipson
Date Submitted:	Group Art Unit Examiner Name	1614
(use as many sheets as necessary)  of 30	Attorney Docket Number	038602-0994

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Cito		U.S. Patent Document		Name of Patentee or Applicant of	Cited Document	Passages or Relevant Figures Appear
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Examiner	Phullic Sailack	Date Considered	1/23/05
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Complete if Known Substitute for form 1449B/PTO **Application Number** INFORMATION DISCLOSURE 12/22/2000 Filing Date STATEMENT BY APPLICANT Ken Lipson First Named Inventor Date Submitted: 1614 Group Art Unit **Examiner Name** (use as many sheets as necessary) 038602-0994 Attorney Docket Number Sheet

<del>,</del>				U.S. PATENT DOCUMENTS	·	
5 in Cito		U.S. Patent		Name of Patentee or Applicant of	Date of Publication of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant
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Examiner Signature	Phullis Souvack	Date Considered	1/23/05
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umber. Substitut	e for form 1449B/PTO	Complete if Known		
INFORMA	ATION DISCLOSURE	Application Number Filing Date	12/22/2000	
STATEM	ENT BY APPLICANT	First Named Inventor	Ken Lipson	
Da	ate Submitted:	Group Art Unit	1614	
(use as mai	ny sheets as necessary)	Examiner Name		
Sheet 5	of 30	Attorney Docket Number	038602-0994	

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		U.S. Patent	Document	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
xaminer nitials*	Cite No. <sup>1</sup>	Number	Kind Code <sup>2</sup> (if known)		MM-DD-YYYY	
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_	miner	Cite No.1	Fore Office <sup>3</sup>	eign Patent Do	cument Kind Code <sup>5</sup> (if known)	Name of Patentee or Applicant of Cited Documents	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T <sup>6</sup>
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1	15 PS	B77	wo	91/13055	A2	FARMITALIA CARLO ERBA SRL	09-05-1991		
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Examiner	Phullis Saivack	Date Considered	1/23/05
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	INFORMAT	ION DISCLO	SURE	Application Number			
	STATEMEN	IT BY APPLI	CANT	Filing Date	12/22/2000		
	Data	Submitted:		First Named Inventor	Ken Lipson		
	Date	Submitteu.		Group Art Unit	1614		
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Sheet	6	of	30	Attorney Docket Number	038602-0994		

				F	OREIGN PATENT DOCUMENT	S		
Examiner	Cite	For	eign Patent D	ocument	Name of Patentee or	Date of Publication of	Pages, Columns, Lines,	T
Initials*	No. <sup>1</sup>	Office <sup>3</sup>	Number <sup>4</sup>	Kind Code <sup>5</sup> (if known)	Applicant of Cited Documents	Cited Document MM-DD-YYYY	Where Relevant Passages or Relevant Figures Appear	T <sup>6</sup>
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	Examiner Signature	Phullis Spirack	Date Considered	1/23/05
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<sup>&</sup>lt;sup>1</sup> Unique citation designation number. <sup>2</sup>See attached Kinds of U.S. Patent Documents. <sup>3</sup>Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). <sup>4</sup>For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. <sup>5</sup>Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. <sup>6</sup>Applicant is to place a check mark here if English language Translation is attached.

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			···	F	OREIGN PATENT DOCUMENT			
Examiner Initials*	Cite No.1	Fore Office <sup>3</sup>	eign Patent D	ocument Kind Code <sup>5</sup> (if known)	Name of Patentee or Applicant of Cited Documents	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	Τ°
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P4	112	wo	99/61422	A1	SUGEN, INC.	12-02-1999		:
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STATEMENT BY APPLICANT

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First Named Inventor Ken Lipson
Group Art Unit 1614

Examiner Name

Attorney Docket Number

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Examiner	Cite	For	eign Patent D	ocument		Name of Patentee or	Date of Publication of	Pages, Columns, Lines,	
Initials*	No.1	Office <sup>3</sup>	Number <sup>4</sup>	Kind Cod (if known		Applicant of Cited Documents	Cited Document MM-DD-YYYY	Where Relevant Passages or Relevant Figures Appear	T⁵
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175	121	DE	2,159,363	1	4	BAYER AG	06-14-1973		х
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F					F	OREIGN PATENT DOCUMENT	S		
	aminer ials*	Cite No.1	For Office <sup>3</sup>	eign Patent Do		Name of Patentee or Applicant of Cited Documents	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T6
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Sheet	10	of	30	Attorney Docket Number	038602-0994	

		OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS	
Exami ner Initials	Cite No.1	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.) date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <sup>6</sup>
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Sheet	11	of	30	Attorney Docket Number	038602-0994			

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B	B208	FOLKMAN and SHING, "Angiogenesis," <u>J. Biol. Chem.</u> 267:10931-10934 (1992) <sup>©</sup> The American Society for Biochemistry and Molecular Biology	
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Date Submitted:		Group Art Unit	1614
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17 of	30	Attorney Docket Number	038602-0994
	Date Submitted: e as many sheets as ned	e as many sheets as necessary)	Date Submitted:  e as many sheets as necessary)  First Named Inventor  Group Art Unit  Examiner Name

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3	B216	GOTTARDIS et al., "Estradiol-Stimulated Growth of MCF-7 Tumors Implanted in Athymic Mice: A Model to Study the Tumoristatic Action of Tamoxifen," <u>J. Steroid Biochem.</u> 30:311-314 (1988) © Pergamon Press	
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Ps	B221	HODGES et al., "Chemical and biological properties of some oxindolidyl-3-methines," <u>Canadian J.</u> <u>Chemistry</u> 46:2189-2194 (1968)	
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	STATEMENT			Filing Date	12/22/2000
				First Named Inventor	Ken Lipson
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				First Named Inventor	Ken Lipson	
	Date	Submitted:		Group Art Unit	1614	
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Sheet	19	of	30	Attorney Docket Number	038602-0994	

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				First Named Inventor	Ken Lipson
	Date	Submitted:		Group Art Unit	1614
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Sheet	20	of	30	Attorney Docket Number	038602-0994

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<b>V</b> 5	B258	MILLAUER et al., "High Affintiy VEGF Binding and Developmental Expression Suggest Flk-1 as a Major Regulator of Vasculogenesis and Angiogenesis," Cell 72:835-846 (1993) © Cell Press	
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P3	B267	NODIFF et al., "Antimalarial Phenanthrene Amino Alcohols. 3. Halogen-containing 9-phenanthrenemethanols," <u>Chemical Abstracts</u> , Vol. 83, abstract no. 188214 (1975)	
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B	B276	QUINN et al., "Fetal liver kinase 1 is a receptor for vascular endothelial growth factor and is selectively expressed in vascular endothelium," <a href="Proc. Natl. Acad. Sci. USA">Proc. Natl. Acad. Sci. USA</a> 90:7533-7537 (1993)				
B	B277	ROZAKIS-ADCOCK et al., "Association of the Shc and Grb2/Sem5 SH2-containing proteins is implicated in activation of the Ras pathway by tyrosine kinases," Nature 360:689-692 (1992)				
B	B278	RUVEDA and GONZALEZ, "Geometric isomerism in benzylideneoxindoles," <u>Spectrochimica Acta</u> 26A:1275-1277 (1970)				
B	B279	RYGAARD and POVLSEN, "Heterotransplantation of a Human Malignant Tumour to 'Nude' Mice," <u>Acta Path. Microbiol. Scand.</u> 77:758-760 (1969)				
Ps	B280	SAINSBURY et al., "Electrochemical Oxidation of Aromatic Ethers. Part 5.1 Further Studies of the Coupling Reactions of Alkoxylated Aralkyl- and Aryl-amides," <u>J.C.S. Perkin I</u> 108-114				
Ps	B281	SAITO and STREULI, "Molecular Characterization of Protein Tyrosine Phosphatases," Cell Growth & Differentation 2:59-65 (1991)   Molecular Biolody Journal of the American Association for Cancer Research				
B	B282	SANDBERG-NORDQVIST et al., "Characterization of Insulin-Like Growth Factor 1 in Human Primary Brain Tumors," Cancer Research 53:2475-2478 (1993)				
PS	B283	SCHINDLER et al., "Über Dibenz[b,f]-azocin-Derivate," Helvetica Chimica Acta 49:985-989 (1966)				
B	B284	SCHLESSINGER and ULLRICH, "Growth Factor Signalling by Receptor Tyrosine Kinases," Neuron 9:383-391 (1992) © Cell Press				

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	Substitute for	form 1449B	I/PTO	Complete if Known		
	INFORMATIC	NDISCLO	SURE	Application Number	08/5/09,842	
STATEMENT BY APPLICANT				Filing Date	12/22/2000	
	Data S	Submittade		First Named Inventor	Ken Lipson	
Date Submitted: (use as many sheets as necessary)				Group Art Unit	1614	
				Examiner Name	Ú. ∩3£∆/2	
Sheet	25	of	30	Attorney Docket Number	038602-0994	

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B	B285	SCHUCHTER et al., "Successful Treatment of Murine Melanoma with Bryostatin 1," Cancer Research 51:682-687 (1991)	,
P5	B286	SEIBERT et al., "Clonal Variation of MCF-7 Breast Cancer Cells in Vitro and in Athymic Nude Mice," Cancer Research 43:2223-2234 (1983)	
75	B287	SHAFIE and GRANTHAM, "Role of Hormones in the Growth and Regression of Human Breast Cancer Cells (MCF-7) Transplanted Into Athymic Nude Mice," <u>J. Natl. Cancer Institute</u> 67:51-56 (1981)	
Ps	B288	SHIBUYA et al., "Nucleotide sequence and expression of a novel human receptor-type tyrosine kinase gene (flt) closely realted to the fms family," Oncogene 5:519-524 (1990)	
B	B289	SHIRAISHI et al., "Specific inhibitors of Tyrosine-Specific Protein Kinase, Synthetic 4-Hydroxycinnamamide Derivatives," <u>Biochemical and Biophysical Research Communications</u> 147:322-328 (1987) <sup>®</sup> Academic Press	
B	B290	SHIRAISHI et al., "Specific Inhibitors of Tyrosine-specific Protein Kinases: Properties of 4-Hydroxycinnamamide Derivatives in Vitro," Cancer Research 49:2374-2378 (1989)	
Ps	B291	SHWEIKI et al., "Vascular endothelial growth factor induced by hypoxia may mediate hypoxia-initiated angiogenesis," Nature 359:843-845 (1992)	•
B	B292	SINGH et al., "Indolinone Derivatives as Potential Antimicrobial Agents," Zentralbl. Mikrobiol. 144:105-109 (1989) copyright VEB Gustav Fischer Verlag Jena	
B	B293	SINGH et al., "Synthesis and Anticonvulsant Activity of New 1-Substituted 1'-Methyl-3-Chloro-2-Oxospiro (Azetidin-3', 4-Indol-2' Ones)," <u>Bollettino Chimico Farmaceutico</u> 133:76-79 (1994)	

Examiner Signature	Y	yllis Spivack	Date Considered	1/23/05

<sup>\*</sup>EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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	INFORMATIO			Application Number	C9/741,0.L	
	STATEMENT BY APPLICANT			Filing Date	12/22/2000	
	Date Submitted:  (use as many sheets as necessary)			First Named Inventor	Ken Lipson	
				Group Art Unit	1614	
				Examiner Name	J. Rsamer	
Sheet	26	of	30	Attorney Docket Number	038602-0994	

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B	B294	SKEHAN et al., "New Colorimetric Cytotoxicity Assay for Anticancer-Drug Screening," <u>J. Natl. Cancer Inst.</u> 82:1107-1112 (1990)	
Ps	B295	SLAMON et al., "Studies of the HER-2/neu Proto-oncogene in Human Breast and Ovarian Cancer," Science 244:707-712 (1989)	
PS	B296	SOLDI et al., "Platelet-Activating Factor (PAF) Induces the Early Tyrosine Phosphorylation of Focal Adhesion Kinase (p125 <sup>FAK</sup> ) in Human Endothelial Cells," <u>Oncogene</u> 13:515-525 (1996) copyright Stockton Press	
Ps	B297	SONGYANG et al., "SH2 Domains Recognize Specific Phosphopeptide Sequences," <u>Cell</u> 72:767-778 (1993) © Cell Press	
3	B298	SONGYANG et al., "Specific Motifs Recognized by the SH2 Domains of Csk, 3BP2, fps/fes, GRB-2, HCP, SHC, Syk and Vav," Molecular and Cellular Biology 14:2777-2785 (1994)   American Society for Microbiology	
B	B299	SPADA, et al., "Small molecule inhibitors of tyrosine kinase activity," Expert Opinion on Therapeutic Patents 5:805-817 (1995) ©Ashley Publications	
PS	B300	STETINOVA et al., "Stereochemistry and Photoisomerisation of Furfurylideneoxindoles," <u>Collection</u> <u>Czechoslov. Chem. Commun.</u> 42:2201-2206 (1977)	
PS	B301	STOLLE, Beilstein Reg. No. 273650, <u>J. Prakt. Chem.</u> , Vol. 2, page 128 (1930)	
B	B302	STOLLE, Beilstein Reg. No. 305045, <u>J. Prakt. Chem.</u> , Vol. 2, page 128 (1930)	

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	INFORMATI			Application Number	25. 71,832		
	STATEMEN	T BY APPL	ICANT	Filing Date	12/22/2000		
	Date	Submitted:	•	First Named Inventor	Ken Lipson		
	Bate	Cabillitiea.		Group Art Unit	1614		
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B	B303	SUMPTER and MILLER, "Chapter IV – Oxindole," in <u>Heterocyclic Compounds With Indole and Carbazole Systems,</u> © Interscience Publishers, Inc., New York, pp. 134-153 (1954)	
B	B304	SUN et al., "Design, Synthesis, and Evaluations of Substituted 3-[(3- or 4-Carboxyethylpyrrol-2-yl) methylidenyl]indolin-2-ones as Inhibitors of VEGF, FGF, and PDGF Receptor Tyrosine Kinases," <u>Journal of Medicinal Chemistry</u> 42: 5120-5130 (1999) ©American Chemical Society	
R	B305	SUN et al, "Synthesis and Biological Evaluations of 3-Substituted Indolin-2-ones: A Novel Class of Tyrosine Kinase Inhibitors That Exhibit Selectivity toward Particular Receptor Tyrosine Kinases," <u>J. Med. Chem.</u> 41:2588-2603 (1998) ©The American Chemical Society	
Ps	B306	SUPERTI-FURGA et al., "A functional screen in yeast for regulators and antagonizers of heterologous protein tyrosine kinases," <u>Nature Biotech.</u> 14:600-605 (1996)	
Ps	B307	SUPERTI-FURGA et al., "Csk inhibition of c-Src activity requires both the SH2 and SH3 domains of Src," EMBO J. 12:2625-2634 (1993) © Oxford University Press	
	B308	TACCONI and MARINONE, "Preparazione e carafferistiche di alcuni 3-ossindolidenderivati," <u>Ricerca Scientifica</u> 38:1239-1244 (1968)	
B	B309	TACCONI et al., "(Z)- and (E)-3-Alkylidene-1,3-dihydroindol-2-ones: Influence of Configuration on the Transmission of the Inductive Effect to the Carbonyl Group," <u>J.C.S. Perkin II</u> 150-154 (1976)	
B	B310	TAKANO et al., "Inhibition of angiogenesis by a novel diaminoanthraquinone that inhibits Protein Kinase C," Mol. Bio. Cell 4:358A at abstract no. 2076 (1993)	
Ps	B311	TERRETT et al., "Combinatorial Synthesis - The Design of Compound Libraries and their Application to Drug Discovery," <u>Tetrahedron</u> 51(30):8135-8173 (1995) copyright Pergamon! all even pages missing!	

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•	INFORMATION	DISCLO	SURE	Application Number	09/741,542	
STATEMENT BY APPLICANT  Date Submitted:				Filing Date	12/22/2000	
				First Named Inventor	Ken Lipson	
	Date Sui	Jiiiileu.		Group Art Unit	1614	
(use as many sheets as necessary)				Examiner Name	J. Reamer	
Sheet	28	of	30	Attorney Docket Number	038602-0994	

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PS	B312	THIO et al., "The Interconversion of 2-(2-Aminophenyl)-3-piperolidinone and 3-(2-piperidylmethyl)-2-indolinone: A Reversible N = N' Transacylation," Notes (1971) 479-482	
B	B313	THOMPSON et al., "Facile Dimerisation of 3-Benzylideneindoline-2-thiones," <u>J. Chem. Soc. Perkin Trans.</u> (I) 1835-1837 (1993)	
PS	B314	TORP et al., "Expression of the Epidermal Growth Factor Receptor Gene in Human Brain Metastases," APMIS 100:713-719 (1992)	
Ps	B315	TRAXLER, "Protein tyrosine kinase inhibitors in cancer treatment," Expert Opinion on Therapeutic Patents 7(6):571-588 (1997) Ashley Publications Ltd.	
P5	B316	TSAI et al., "The Effect of 3,3-Di-Pyridyl Methyl-1-Phenyl-2-Indolinone on the Nerve Terminal Currents of Mouse Skeletal Muscles," Neuorpharmacology 31:943-947 (1992) Pergamon Press	
PS	B317	TUZI et al., "Expression of growth factor receptors in human brain tumours," <u>Br. J. Cancer</u> 63:227-233 (1991)	
Ps	B318	TWAMLEY-STEIN et al., "The Src family tyrosine kinases are required for platelet-derived growth factor-mediated signal transduction in NIH 3T3 cells," <u>Proc. Natl. Acad. Sci. USA</u> 90:7696-7700 (1993)	
Ps	B319	ULLRICH and SCHLESSINGER, "Signal Transduction by Receptors with Tyrosine Kinase Activity," <u>Cell</u> 61:203-212 (1990) copyright Cell Press	
PS	B320	VAISMAN et al., "Characterization of the Receptors for Vascular Endothelial Growth Factor," <u>J. Biol. Chem.</u> 265:19461-19466 (1990) © The American Society for Biochemistry and Molecular Biology	

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	Date	Submitted:		First Named Inventor	Ken Lipson			
	Date	Submitted.		Group Art Unit	1614			
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Sheet	29	of	30	Attorney Docket Number	038602-0994			

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B	B321	VARMA and GUPTA, "Nucleophilic Reactions of 2-Methyl-3-(4'-carbomethoxyphenyl)-4-quinazolinones with 2-Indolinones," <u>J. Indian Chem. Soc.</u> 66:804-805 (1989) © The Indian Chemical Society	
Ps	B322	VOLLER et al., "Ch. 45 – Enzyme-Linked Immunosorbent Assay," in Manual of Clinical Immunology, 2 <sup>nd</sup> edition, Rose and Friedman editors, American Society of Microbiology, Washington, D.C., pp. 359-371 (1980); @ American Society for Microbiology	
PS	B323	WAHL et al., "3-benzilidene-5-methyl-1,3-dihydroindol-2-one," <u>Ann. Chim.</u> 350 (1926), DATABASE CROSSFIRE, Beilstein Reference No. 2-21-00-00290	
PS	B324	WAHL et al., "Chimie Organique - Sur les iso-indogenides," <u>C.R. Hebd. Seances Acad. Sci.</u> 149:132-134 (1909)	
B	B325	WAHL, Beilstein Reg. No. 191439, <u>Bull. Soc. Chim. Fr.</u> , page 1038 (1909)	
B	B326	WAHL, Beilstein Reg. No. 231732, <u>Bull. Soc. Chim. Fr.</u> , pages 1035-1038 (1909)	
PS	B327	WALKER, "Synthesis of a α-(p-Aminophenyl)- and α-(p-Chlorophenyl)-β-aryl-propionitriles by Catalytic Reduction of Stilbenenitriles," <u>J. Med. Chem.</u> 8:583-588 (1965)	
RS	B328	WALKER et al., "Synthesis of New 3-(Pyridylmethylene)-, 3-(Pyridylmethyl)-, 3-(Piperidylmethyl)-, and 3-(β-Alkylaminoethyl)-2-indolinones. The Reduction of Isoindogenides, Nitro Compounds, and Pyridines in a Series of 2-Indolinones," <u>J. Med. Chem.</u> 8:626-637 (1965)	
Ps	B329	WARRI et al., "Estrogen Suppression of erbB2 Expression is Associated with Increased Growth Rate of ZR-75-I Human Breast Cancer Cells <u>In Vitro</u> and in Nude Mice," <u>Int. J. Cancer</u> 49:616-623 (1991) © Wiley-Leiss, Inc.	

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	Date	Submitted:		First Named Inventor	Ken Lipson		
	Date	Submitted.		Group Art Unit	1614		
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1/5	B330	WEIDNER et al., "Tumor Angiogenesis and Metastasis Correlation in Invasive Breast Carcinoma," New England J. Medicine 324:1-7 (1991)   Massachusetts Medical Society				
B	B331	WINKELMANN et al., "Chemotherapeutically Active Nitro Compounds: 4. 5-Nitroimidazoles (Part I)," ArzneimForsch./Drug Res. 27:2251-2263 (1977)				
B	B332	WRIGHT et al., "Cyclic Hydroxamic Acids Derived from Indole," <u>J. Am. Chem. Soc.</u> 78:221-224 (1956)				
PS	B333	WRIGHT et al., "Inhibition of Angiogenesis in Vitro and In Ovo With an Inhibitor of Cellular Protein Kinases, MDL 27032," <u>J. Cellular Physiology</u> 152:448-457 (1992)				
PS	B334	YOUNG and BABBITT, "2-(2-Methyl-3-indolyl)-1,4-benzoquinone, a Reversible Redox Substrate at the Carbon-Paste Electrode in Acidic Aqueous-Ethanolic Media," <u>J. Org. Chem.</u> 47:1571-1572 (1982) copyright Am. Chem. Soc.				
PS	B335	ZAMAN et al., "Tyrosine Kinase Activity of Purified Recombinant Cytoplasmic Domain of Platelet-Derived Growth Factor β-Receptor (β-PDGFR) and Discovery of a Novel Inhibitor of Receptor Tyrosine Kinases," Biochemical Pharmacology 57:57-64 (1999) ©Elsevier Science Inc.				
PS	B336	ZHANG et al., "Microtubule Effects of Welwistatin, a Cyanobacterial Indolinone that Circumvents Multiple Drug Resistance," Molecular Pharmacology 49:228-234 (1996) The American Society for Pharmacology and Experimental Pharmaceutics				
PS	B337	ZHUNGIETU et al., "Reaction of Indoles and 2-Ketoindolines With Some Aldehydes," <u>Chemical Abstracts</u> , Vol. 78, abstract no. 111201 (1990)				

Examiner Signature	Phyllis Sawack	Date Considered	1/23/05

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